6. Tablet in accordance with Claim 1 wherein the quantity of lubricating agent is in the range 0.2 to 10 parts per 1000 (weight of lubricating agent / total weight of tablet), and is preferably in the range 3 to 6 parts per 1000 (weight of lubricating agent / total weight of tablet).

7. Tablet in accordance with Claim 1 wherein the lubricating agent has a particle size distribution such that its constituent particles adhere when it is sprayed against a surface, preferably less than 30 microns and more preferably still, less than 10 microns.

8. Tablet in accordance with Claim 1 wherein the disintegrating agent is selected from the group including in particular cross-linked sodium carboxymethylcellulose, known in the industry as croscarmellose, crospovidone and their mixtures.

9. Tablet in accordance with Claim 1 wherein the mixture of excipients may include a permeabilising agent, a solubilising agent, sweeteners, flavors and colorings.

10. Tablet in accordance with Claim 1 wherein said tablet is designed to be packaged in blisters composed entirely of aluminum, which may in addition include a cover of a plastic material which is to be torn off before opening.

11. Process for the production of a tablet in accordance with Claim 1 wherein the process involves the following sequence of steps:

choosing, firstly, an active substance in the form of coated microcrystals or microgranules, and secondly, a set of excipients including a disintegrating agent, a soluble agent, and also a lubricating agent;

mixing the active substance and the excipients with the exception of at least the greater part of the lubricating agent;

feeding a quantity of this mixture necessary to form a tablet into the cavity of a compression device within which